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Notice of Allowability	Application No.	Applicant(s)	
	09/711,782	BITNER ET AL.	
	Examiner Jezia Riley	Art Unit 1637	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address--

All claims being allowable, PROSECUTION ON THE MERITS IS (OR REMAINS) CLOSED in this application. If not included herewith (or previously mailed), a Notice of Allowance (PTOL-85) or other appropriate communication will be mailed in due course. **THIS NOTICE OF ALLOWABILITY IS NOT A GRANT OF PATENT RIGHTS.** This application is subject to withdrawal from issue at the initiative of the Office or upon petition by the applicant. See 37 CFR 1.313 and MPEP 1308.

1. This communication is responsive to Amdt filed 9/29/03.
2. The allowed claim(s) is/are 1-51.
3. The drawings filed on _____ are accepted by the Examiner.
4. Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
 - a) All b) Some* c) None of the:
 1. Certified copies of the priority documents have been received.
 2. Certified copies of the priority documents have been received in Application No. _____.
 3. Copies of the certified copies of the priority documents have been received in this national stage application from the International Bureau (PCT Rule 17.2(a)).

* Certified copies not received: _____.

5. Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).
 - (a) The translation of the foreign language provisional application has been received.
6. Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.

Applicant has THREE MONTHS FROM THE "MAILING DATE" of this communication to file a reply complying with the requirements noted below. Failure to timely comply will result in ABANDONMENT of this application. **THIS THREE-MONTH PERIOD IS NOT EXTENDABLE.**

7. A SUBSTITUTE OATH OR DECLARATION must be submitted. Note the attached EXAMINER'S AMENDMENT or NOTICE OF INFORMAL PATENT APPLICATION (PTO-152) which gives reason(s) why the oath or declaration is deficient.
8. CORRECTED DRAWINGS must be submitted.
 - (a) including changes required by the Notice of Draftsperson's Patent Drawing Review (PTO-948) attached
 - 1) hereto or 2) to Paper No. _____.
 - (b) including changes required by the proposed drawing correction filed _____, which has been approved by the Examiner.
 - (c) including changes required by the attached Examiner's Amendment / Comment or in the Office action of Paper No. 11/03.

Identifying indicia such as the application number (see 37 CFR 1.84(c)) should be written on the drawings in the front (not the back) of each sheet.

9. DEPOSIT OF and/or INFORMATION about the deposit of BIOLOGICAL MATERIAL must be submitted. Note the attached Examiner's comment regarding REQUIREMENT FOR THE DEPOSIT OF BIOLOGICAL MATERIAL.

Attachment(s)

1 <input type="checkbox"/> Notice of References Cited (PTO-892)	2 <input type="checkbox"/> Notice of Informal Patent Application (PTO-152)
3 <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948)	4 <input type="checkbox"/> Interview Summary (PTO-413), Paper No. _____.
5 <input checked="" type="checkbox"/> Information Disclosure Statements (PTO-1449), Paper No. <u>11/03</u>	6 <input checked="" type="checkbox"/> Examiner's Amendment/Comment
7 <input type="checkbox"/> Examiner's Comment Regarding Requirement for Deposit of Biological Material	8 <input type="checkbox"/> Examiner's Statement of Reasons for Allowance
	9 <input type="checkbox"/> Other _____.

EXAMINER'S AMENDMENT

An examiner's amendment to the record appears below. Should the changes and/or additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be submitted no later than the payment of the issue fee.

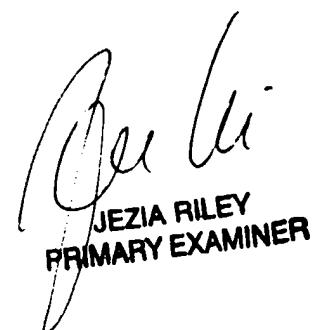
The formal Drawings are required.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jezia Riley whose telephone number is 703-305-6855. The examiner can normally be reached on 9:30AM - 5:00PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Benzion can be reached on 703-308-1119. The fax phone number for the organization where this application or proceeding is assigned is (703) 872-9306.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 703-308-0196.

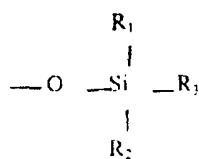
Wednesday, November 12, 2003



JEZIA RILEY
PRIMARY EXAMINER

ALLOWED CLAIMS/ TJ

1. (Currently amended) A method of clearing a solution of disrupted biological material, according to steps comprising:
 - (a) providing a first silanized silica matrix, comprising a silica solid phase with a plurality of silane ligands covalently attached thereto, wherein each of the plurality of ligands has a neutral charge in a first solution; and
 - (b) combining the first silanized silica matrix with the first solution, comprising a disrupted biological material, a target nucleic acid material, and a chaotropic salt at a concentration sufficient to promote selective adsorption of the disrupted biological material to the matrix, thereby forming a first complex between the silanized silica matrix and the disrupted biological material.
2. (Original) The method of claim 1, wherein the disrupted biological material is a bacterial cell lysate.
3. (Original) The method of claim 1, wherein the disrupted biological material is disrupted plant matter.
4. (Original) The method of claim 1, wherein the chaotropic salt concentration in step (b) is at least about 0.5 M.
5. (Original) The method of claim 1, wherein the each ligand in the plurality of silane ligands is of the general formula:



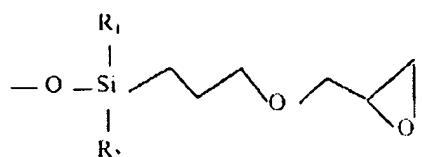
wherein R_1 and R_2 are each a subunit selected from the group consisting of a hydrocarbon chain having from 1 to 5 carbon atoms, an alkoxy having from 1 to 5 carbon atoms, a halogen atom, a hydrogen atom, a hydroxy, an alkyl chain having from 4 to 10

Art Unit: 1637

carbon atoms interrupted by an oxy residue wherein up to five of the carbon atoms is substituted by a group selected from the group consisting of a halogen, an alkoxy having from 1 to 3 carbon atoms, a cyano having from 1 to 3 carbon atoms, a hydroxy, and a linkage to another silane ligand; and

wherein R₃ is a hydrocarbon chain having from 1 to 20 carbon atoms substituted by at least one hydroxy, an alkyl chain having from 4 to 20 carbon atoms interrupted by at least one oxy group wherein up to ten carbon atoms are replaced by a moiety selected from the group consisting of a halogen, a cyano having from 1 to 3 carbon atoms, an alkoxy having from 1 to 3 carbon atoms, a hydroxy, and an epoxy

6. (Previously presented) The method of claim 1, wherein each ligand in the plurality of silane ligands is of the general formula:



wherein, R₁ and R₂ are each independently selected from the group consisting of -OH, -CH₃, -OCH₃, and a linkage to another silane ligand to generate a higher order polymer.

7. (Original) The method of claim 1, wherein the silica solid phase is a first silica magnetic particle.

8. (Original) The method of claim 1, further comprising a step of separating the first complex from the first solution, thereby producing a cleared solution.

9. (Original) The method of claim 8, further comprising a step of combining the cleared solution with a second silica matrix in a second solution, wherein the target nucleic acid specifically adsorbs to the second silica matrix, thereby forming a second complex.

10. (Original) The method of claim 9, wherein the second silica matrix comprises a plurality of second silica magnetic particles.

11. The method of claim 9, wherein the second silica matrix is a plurality of second silanized silica magnetic particles, and the second solution has a pH of up to about 8.0.

12. (Original) The method of claim 9, wherein the first silica matrix and the second silica matrix are the same.

13. (Currently amended) A method of clearing a solution of disrupted biological material, according to steps comprising:

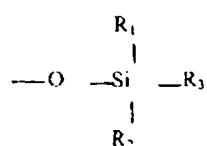
- (a) providing a first silanized silica magnetic particle comprising a silica magnetic particle with a plurality of silane ligands covalently attached thereto;
- (b) combining the first silanized silica magnetic particle with a first solution, comprising a disrupted biological material, a target nucleic acid, and a chaotropic salt concentration sufficiently high to promote selective adsorption of the disrupted biological material to the silanized silica magnetic particle, thereby forming a first complex between the silanized silica matrix and the disrupted biological material;
- (c) separating the first complex from the first solution, thereby forming a cleared solution.

14. (Original) The method of claim 13 wherein the disrupted biological material is a bacterial cell lysate.

15. (Original) The method of claim 13 wherein the disrupted biological material is disrupted plant matter.

16. (Original) The method of claim 13, wherein the first solution further comprises a chaotropic salt at a concentration of up to about 3.5M.

17. (Original) The method of claim 13, wherein the each of the plurality of silane ligands is of the general formula:



wherein R₁ and R₂ are each a subunit selected from the group consisting of a hydrocarbon chain having from 1 to 5 carbon atoms, an alkoxy having from 1 to 5 carbon atoms, a halogen atom, a hydrogen atom, a hydroxy, an alkyl chain having from 4 to 10 carbon atoms interrupted by an oxy residue wherein up to five of the carbon atoms is substituted by a group selected from the group consisting of a halogen, an alkoxy having from 1 to 3 carbon atoms, a cyano having from 1 to 3 carbon atoms, and a hydroxy; wherein R₃ is a hydrocarbon chain having from 1 to 20 carbon atoms substituted by at least one hydroxy, an alkyl chain having from 4 to 20 carbon atoms interrupted by at least one oxy group wherein up to ten carbon atoms are replaced by a moiety selected from the group consisting of a halogen, a cyano having from 1 to 3 carbon atoms, an alkoxy having from 1 to 3 carbon atoms, a hydroxy, an epoxy, and a linkage to another silane ligand.

18. (Original) The method of claim 13, wherein the first complex is separated from the first solution in the presence of a magnetic field.

19. (Original) The method of claim 13, wherein the first complex is separated from the first solution by centrifugation.

20. (Original) The method of claim 13, further comprising a step of combining the cleared solution with a second silica matrix in a second solution, wherein the target nucleic acid specifically adsorbs to the second silica matrix, thereby forming a second complex.

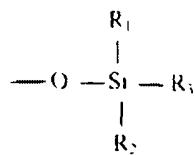
21. (Original) The method of claim 20, wherein the second silica matrix is a second silanized silica magnetic particle comprising a silica magnetic particle solid phase with a plurality of silane ligands covalently attached thereto.

22. (Original) The method of claim 21, wherein the first silanized silica magnetic particle and the second silanized silica magnetic particle are the same.

23. (Original) The method of claim 20, wherein the second silica matrix is a silica magnetic particle.

24. (Original) A method of isolating a target nucleic acid from a nucleic acid adsorption solution, comprising the steps of:

(a) providing a silanized silica matrix comprising a silica solid phase with a plurality of silane ligands covalently attached thereto, wherein each of the plurality of silane ligands is of the general formula:



wherein R_1 and R_2 are each a subunit selected from the group consisting of a hydrocarbon chain having from 1 to 5 carbon atoms, an alkoxy having from 1 to 5 carbon atoms, a halogen atom, a hydrogen atom, a hydroxy, an alkyl chain having from 4 to 10 carbon atoms interrupted by an oxy residue wherein up to five of the carbon atoms is substituted by a group selected from the group consisting of a halogen, an alkoxy having from 1 to 3 carbon atoms, a cyano having from 1 to 3 carbon atoms, and a hydroxy; wherein R_3 is a hydrocarbon chain having from 1 to 20 carbon atoms substituted by at least one hydroxy, an alkyl chain having from 4 to 20 carbon atoms interrupted by at least one oxy group wherein up to ten carbon atoms are replaced by a moiety selected from the group consisting of a halogen, a cyano having from 1 to 3 carbon atoms, an alkoxy having from 1 to 3 carbon atoms, a hydroxyl, an epoxy, and a linkage to another silane ligand;

- (b) combining the silanized silica matrix with a nucleic acid adsorption solution having a pH of up to about pH 8.0, the nucleic acid adsorption solution comprising the target nucleic acid and at least one non-target material, wherein the target nucleic acid selectively adsorbs to the silanized silica matrix, thereby forming a complex; and
- (c) separating the complex from the nucleic acid adsorption solution.

25. (Original) The method of claim 24, wherein the nucleic acid adsorption solution comprises a vegetable oil.

26. (Original) The method of claim 24, wherein the nucleic acid adsorption solution further comprises a concentration of low molecular weight alcohol sufficient to promote adsorption of the target nucleic acid to the second silanized silica matrix.

27. (Original) The method of claim 24, wherein the adsorption solution further comprises 0.2M to 1.2M of a chaotropic salt.

28. (Original) The method of claim 27, wherein the chaotropic salt is selected from the group consisting of guanidine hydrochloride and guanidine thiocyanate.

29. (Original) The method of claim 24, wherein the silica solid phase of the silica matrix is a silica magnetic particle.

30. (Original) The method of claim 29, wherein the complex is separated from the nucleic acid adsorption solution in the presence of a magnetic field.

31. (Original) The method of claim 24, further comprising washing the complex in a wash solution having a pH of up to about 8.0.

32. (Previously presented) The method of claim 31, wherein the wash solution comprises a concentration of at least about 30% of a low molecular weight alcohol.

33. (Original) The method of claim 24, further comprising combining the complex with an elution solution having a pH of at least about 8.0, thereby desorbing the target nucleic acid from the complex.

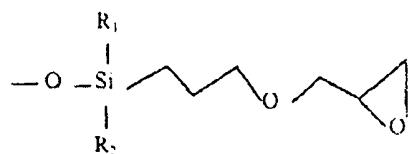
34. (Original) The method of claim 33, wherein the elution solution is a buffer having a pH of at least about 9.0.

35. (Original) The method of claim 24, wherein the target nucleic acid is selected from the group consisting of plasmid DNA, genomic DNA, and total RNA.

36. (Original) The method of claim 24, wherein the target nucleic acid is double-stranded linear DNA with a molecular weight of at least about 25 base pairs and up to about 60 kilobase pairs.

37. (Previously presented) A method of isolating a target nucleic acid from a nucleic acid adsorption solution using a silanized silica magnetic particle, comprising the steps of:

- (a) providing a silanized silica magnetic particle, comprising a silica magnetic particle with a plurality of silane ligands covalently attached thereto, wherein each of the plurality of silane ligands is of a general formula:



wherein, in each formula, R₁ and R₂ are each independently selected from the group consisting of -OH, -CH₃, -OCH₃, or -OCH₂CH₃, and a linkage to another silane ligand to generate a higher order polymer;

- (b) combining the silanized silica magnetic particle with a nucleic acid adsorption solution having a pH of up to about pH 8.0, the nucleic acid adsorption solution comprising the target nucleic acid and at least one non-target material, wherein the target nucleic acid selectively adsorbs to the silanized silica magnetic particle, thereby forming a complex; and
- (c) separating the complex from the adsorption solution.

38. (Original) The method of claim 37, wherein the adsorption solution has a pH of up to about 8.0.

39. (Original) The method of claim 37, wherein the adsorption solution comprises a vegetable oil.

40. (Original) The method of claim 37, wherein the adsorption solution comprises the target nucleic acid from an agarose gel slice and the agarose gel.

41. (Original) The method of claim 37, wherein the adsorption solution further comprises a concentration of low molecular weight alcohol sufficient to promote adsorption of the target nucleic acid to the silanized silica magnetic particle.

42. (Original) The method of claim 37, wherein the adsorption solution further comprises a chaotropic salt.

43. (Original) The method of claim 42, wherein the chaotropic salt is selected from the group consisting of guanidine hydrochloride and guanidine thiocyanate.

44. (Original) The method of claim 37, further comprising washing the complex in a wash solution having a pH of up to about 8.0.

45. (Original) The method of claim 44, wherein the wash solution comprises a concentration of at least about 30% of a low molecular weight alcohol.

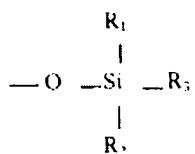
46. (Original) The method of claim 37, further comprising combining the complex with an elution solution having a pH of at least about 8.0, thereby eluting the target nucleic acid from the complex.

47. (Original) The method of claim 37, wherein the target nucleic acid is selected from the group consisting of plasmid DNA, genomic DNA, and total RNA.

48. (Original) The method of claim 37, wherein the target nucleic acid is DNA with a molecular weight of at least 25 base pairs and up to about 60 kilobase pairs.

49. (Original) A kit comprising, in a single container:

a plurality of silanized silica magnetic particles comprising a silica solid phase with at least one silane ligand covalently attached to the surface of each particle, the silane ligand having a structure of formula:



wherein R_1 and R_2 are each a subunit selected from the group consisting of a hydrocarbon chain having from 1 to 5 carbon atoms, an alkoxy having from 1 to 5 carbon atoms, a halogen atom, a hydrogen atom, a hydroxy, an alkyl chain having from 4 to 10 carbon atoms interrupted by an oxy residue wherein up to five of the

carbon atoms is substituted by a group selected from the group consisting of a halogen, an alkoxy having from 1 to 3 carbon atoms, a cyano having from 1 to 3 carbon atoms, and a hydroxy; wherein R_3 is a hydrocarbon chain having from 1 to 20 carbon atoms substituted by at least one hydroxy, an alkyl chain having from 4 to 20 carbon atoms interrupted by at least one oxy group wherein up to ten carbon atoms are replaced by a moiety selected from the group consisting of a halogen, a cyano having from 1 to 3 carbon atoms, an alkoxy having from 1 to 3 carbon atoms, a hydroxy, an epoxy, and a linkage to another silane ligand.

50. (Previously presented) The method of claim 31, wherein the wash solution is water.

51. (Previously presented) The method of claim 44, wherein the wash solution is water.